camovis.

CLINICAL OPERATIONS

Your efficient path to enhanced study execution

Visit us: camovis.de

Challenges

Many trial centers are under pressure: staff shortages, high demands and limited resources make it difficult to conduct clinical trials in a structured and in the expected high-quality manner. The result: delays, team overload, and risks to data quality and patient care.

- **Staffing of trial sites**
  Insufficient availability of qualified personnel to fully meet the requirements in conducting clinical trials.
- **Quality and speed**
  Resource shortages reduce the quality of care for study participants, prolong studies, and increase error risks.
- **Workload of the study team**
  Employee overload leads to stress, errors and staff turnover which slows down overall study performance.
- **On-site and remote patient care**
  Barriers to accessing patient populations and gaps in care can resut to low recruitment rates, deviations from the protocol, and incomplete data.

Why us

- ✓ Hands-on expertise
- ✓ Proactive project planning
- ✓ Efficient site management
- ✓ Well-structured workflows
- ✓ Smooth communication
- ✓ Strict ICH/GCP fulfillment

We care

Camovis Clinical Service performs essential tasks in the implementation and execution of study procedures including patient care in clinical trials. With our experienced clinical research professionals we provide a flexible patient-focused service — at the trial site, at the patient's home or remotely.

Our employees provide support during study visits: vital sign measurements, ECGs, blood draws, IMP handling and blood sample logistics. They assist with the use of digital tools such as eDiaries and wearables, ensuring valid and complete data collection. This support seamlessly complements your staff, filling gaps or temporarily taking over entire roles in hybrid and decentralized study settings.

Whether as a long-term addition or to bridge staffing shortages, we support our customers where it is needed. Goal-oriented, sustainable, and efficient.

camovis.

CLINICAL OPERATIONS

Your efficient path to enhanced study execution

Visit us: camovis.de

Service Packages

Preparations & execution of study visits

- Preparation, support, and follow-up of study visits
- IMP handling and preparation (blinded and unblinded)
- Patient support and training on the administration of study medication and the use of medical devices
- Collection and collation of clinical source data

Examinations & interventions

- Measurement of vital signs (RR, HR, Temp., resp., O2)
- Venous punctures and blood sampling
- Performance of functional tests (e.g., ECG, spirometry, 6-minute walk test)
- Surveillance of IMP application (incl. infusion therapies) and medical devices

Sample handling & laboratory logistics

- Collection, processing, and storage of blood samples, urine samples, and tissue samples
- Centrifugation, aliquoting and documentation of samples
- Prepare and ship samples in accordance with the IATA regulations
- Communication with lab-and logistic providers

Decentralized Trials (DCT) - Clinical Operations

- Camovis Clinical Operations Services support the decentralized conduct of clinical trials. Please refer to the DCT Service Information to find out more about the wide range of available options.

Let`s get to work

.

Camovis GmbH

E-Mail: info@camovis.de
Phone: +49 (0)30 / 233 299 96

Feel free to contact us — we're here and happy to help! :)

WE MAKE YOUR RESEARCH FLY